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(FILE 'HOME' ENTERED AT 09:55:08 ON 10 MAY 2004)

7 S L10 AND DELBRUECKII

2 DUP REMOVE L11 (5 DUPLICATES REMOVED)

L1 L2 L3	FILE	'MEDLINE, CAPLUS, BIOSIS' ENTERED AT 09:55:29 ON 10 MAY 2004 3166 S LACTOBACILLUS AND DELBRUECKII 64 S L1 AND (MODIFICATION OR MODIFY OR TRANSFORM OR INTEGRATE) 30 S L2 AND (GENE OR GENETIC OR CHROMOSOME)
пэ		50 b Hz find (Child On Children)
	FILE 2004	'MEDLINE, CAPLUS, BIOSIS, SCISEARCH' ENTERED AT 09:58:34 ON 10 MAY
L4		43 S L3
L5		3 S L4 AND REVIEW
L6		2 S L4 AND (DIFFICULT OR CHALLENGE OR OBSTACLE OR DIFFICULTIES O
L7		19 S GENETIC MODIFICATION AND LACTOBACILLI
L8		11 DUP REMOVE L7 (8 DUPLICATES REMOVED)
L9		39616 S CHROMOSOME AND (INSERTION OR INSERT OR INTEGRATE OR INTEGRATI
L10		22 S L9 AND LACTOBACILLI

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L11

L12

WO 2000-FR2565 W 20000915

AB The invention concerns a method for modifying the chromosomal genetic information of Lactobacillus delbrueckii, using a conditional integrator plasmid. The plasmid contains a  $\theta$  replication system of plasmid pIP501 or a related plasmid. Plasmid pIP501 is stable and can replicate in L. delbrueckii at 35-37° but cannot replicate and is unstable at 42°. The method for modifying the L. delbrueckii chromosome comprises (a) insertion of a DNA sequence capable of integrating into the bacterial chromosome and a selectable marker into the thermosensitive integrator plasmid, (b) introduction of this plasmid into the bacteria and multiplication of the transformants under conditions favoring plasmid replication and maintenance, (c) multiplication of selectable marker-expressing bacteria under conditions nonpermissive for plasmid replication and stability, and, optionally, (d) recovery of bacteria expressing the selectable marker from the previous step. Sequences capable of inserting into the bacterial chromosome include transposons and insertion sequences such as IS1223 and IS1201. The invention also concerns integrator plasmids for use in implementing said method. Plasmids of the invention include pVI49, containing IS1223, and pVI52, containing IS1201.

ANSWER 3 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN 1.2

2001:229062 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER:

INVENTOR (S):

134:263403

TITLE:

Lactobacillus delbrueckii strain, its use

for screening plasmids, and plasmids so obtained Serror, Pascale; Fremaux, Christophe; Benbadis,

Laurent; Maguin, Emmanuelle

PATENT ASSIGNEE(S):

Institut National de la Recherche Agronomique (INRA),

Fr.; Compagnie Gervais Danone; Rhodia Food

SOURCE:

PCT Int. Appl., 30 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

French

13

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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KIND DATE
                                  APPLICATION NO. DATE
PATENT NO.
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               A1
                     20010329
                                  WO 2000-FR2564 20000915
WO 2001021818
   W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
       CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
       HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
       LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
       SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,
       YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
   RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
       DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
       CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
FR 2798669
                     20010323
                                  FR 1999-11683
                A1
                                                    19990917
                     20040220
FR 2798669
                B1
EP 1216305
                A1
                     20020626
                                   EP 2000-964296
                                                    20000915
   R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, MC, IE, SI,
       LT, LV, FI, RO, MK, CY, AL
                                 FR 1999-11683
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PRIORITY APPLN. INFO.:

A 19990917 WO 2000-FR2564 W 20000915

The invention concerns a novel strain of Lactobacillus delbrueckii called VI104 which may be used to screen for plasmids useful for transformation of Lactobacillus. Preferred plasmids identified with this strain include pJK650, pGB305 $\Delta$ , and pLEM415.

REFERENCE COUNT:

THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L5 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1996:239443 CAPLUS

DOCUMENT NUMBER:

124:308582

TITLE:

Molecular tools for the genetic modification of dairy lactobacilli

AUTHOR (S):

Klein, Juergen R.; Ulrich, Christof; Wegmann, Udo; Meyer-Barton, Elke; Plapp, Roland; Henrich, Bernhard

CORPORATE SOURCE:

Fachbereich Biologie, Universitat Kaiserslautern,

Kaiserslautern, D-67653, Germany

SOURCE:

Systematic and Applied Microbiology (1996), 18(4),

493-503 CODEN: S Fischer

CODEN: SAMIDF; ISSN: 0723-2020

PUBLISHER:

DOCUMENT TYPE:

Journal: General Review

LANGUAGE:

GE: English

A review and discussion with 52 refs. The possibility that starter strains of lactic acid bacteria may be deliberately endowed with desired properties by the use of mol. techniques is an attractive perspective for the dairy industry. To contribute to the development of appropriate genetic modification systems for lactobacilli, we tested the accessibility of these bacteria to electrotransformation, and we designed a strategy for the construction and use of food-grade plasmid vectors. Successful electroporation was achieved with two dairy strains of Lactobacillus casei and Lb. delbrueckii ssp. lactis. Aiming at the construction of vectors, replicons of cryptic plasmids from various starter lactobacilli were identified and characterized on the mol. and functional levels. provide the resulting vectors with food-grade markers, a number of peptidase and transport genes of the proteolytic system of Lb. delbrueckii ssp. lactis DSM7290 were isolated and sequenced, and features of the resp. gene products were investigated. Over expression of two of these peptidase genes in Lactobacillus had no effects on cell growth, but altered the composition of the growth medium.

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